CLARITIN- loratadine tablet
Bayer HealthCare LLC.

Non-Drowsy*Claritin ®
loratadine tablets 10 mg/antihistamine

Drug Facts

Active ingredient (in each tablet)

Loratadine 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- itching of the nose or throat

Warnings

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours

children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information (Blister foil units)

- safety sealed: do not use if the individual blister unit imprinted with Claritin ® is open or torn
- store between 20° to 25°C (68° to 77°F)
- protect from excessive moisture

Other information (Bottles)

- Tamper-evident: do not use if foil seal under cap, printed with "SEALED for YOUR PROTECTION" is missing, open or broken
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate

Questions or comments?

1-800-CLARITIN (1-800-252-7484) or **www.claritin.com**

PRINCIPAL DISPLAY PANEL - 10 Tablet Carton

Original Prescription

Strength

Non-Drowsy*

Claritin®

loratadine tablets 10 mg/antihistamine

24 Hour

Relief of:

- *Sneezing*
- *Runny Nose*
- *Itchy, Watery Eyes*
- *Itchy Throat or Nose*

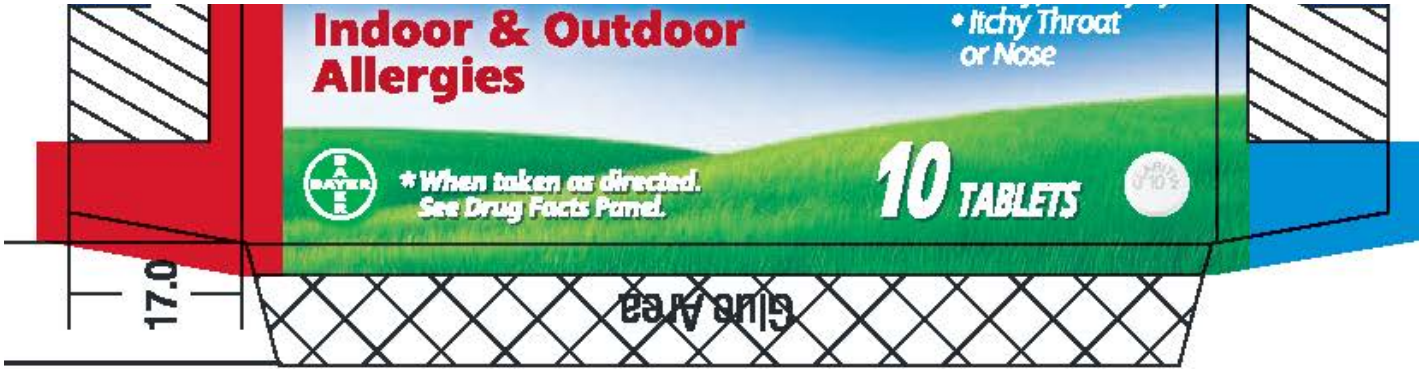
Indoor & Outdoor

Allergies

***When taken as directed.**

See Drug Facts Panel.

10 Tablets



PRINCIPAL DISPLAY PANEL - 45 Count Window Box

Original Prescription Strength

Non-Drowsy *

Claritin ®

loratadine tablets 10 mg/antihistamine

24 Hour

Relief of:

Sneezing
Runny Nose
Itchy, Watery Eyes
Itchy Throat or Nose

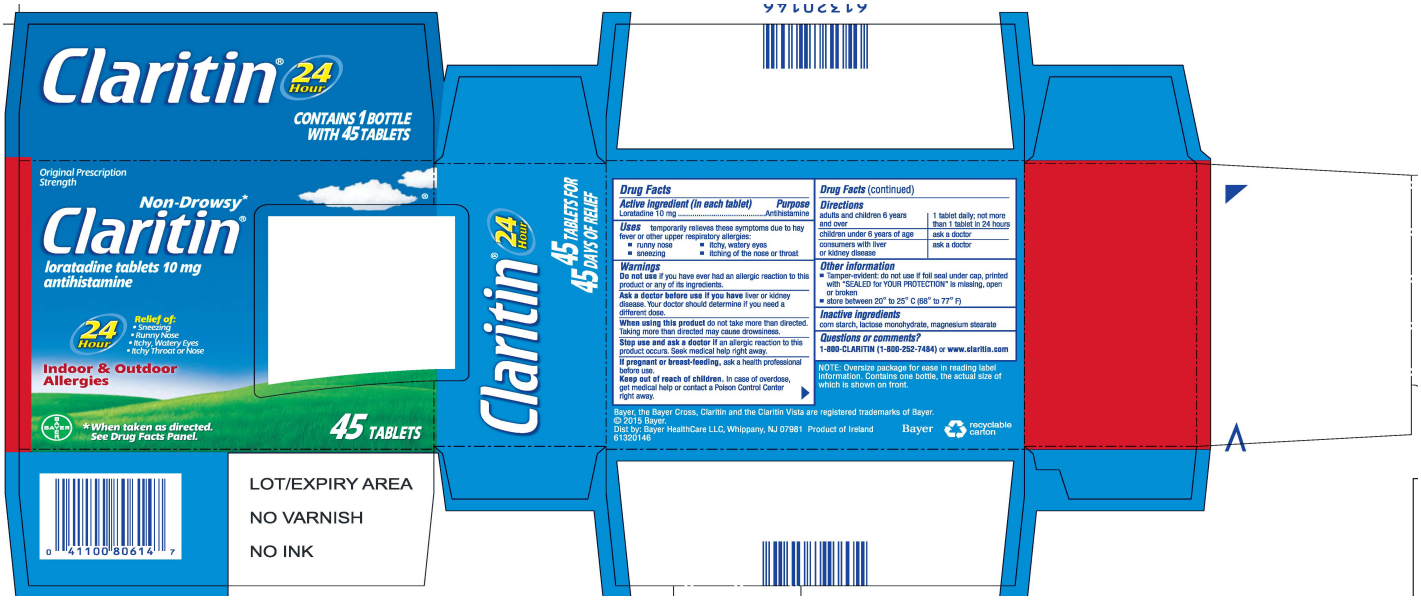
Indoor & Outdoor

Allergies

* When taken as directed.

See Drug Facts Panel.

45 TABLETS



PRINCIPAL DISPLAY PANEL - 50 Tablet Carton

Non-Drowsy*

Claritin®

loratadine tablets 10 mg/antihistamine

24 Hour

Relief of:

- *Sneezing*
- *Runny Nose*
- *Itchy, Watery Eyes*
- *Itchy Throat or Nose*

Indoor & Outdoor

Allergies

***When taken as directed.**

See Drug Facts Panel.

50 1-Count Pouches

Total 50 Tablets


Non-Drowsy*
Claritin®
24 Hour
loratadine tablets 10 mg/antihistamine
Relief of:
• Sneezing
• Runny Nose
• Itchy, Watery Eyes
• Itchy Throat or Nose
Indoor & Outdoor Allergies
*When taken as directed. See Drug Facts Panel.
50 1-Count Pouches
Total 50 Tablets
15 1/8


Claritin®
24

Claritin 24 Hour

Drug Facts

Active ingredient (in each tablet) — **Purpose**

Loratadine 10 mg ..... Antihistamine

Uses temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

Warnings

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Drug Facts (continued)

Directions

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

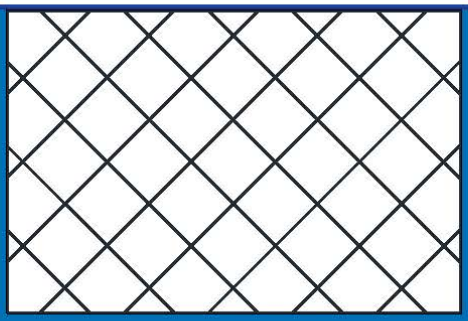
- safety sealed: do not use if the individual blister unit imprinted with Claritin® is open or torn
- store between 20° to 25°C (68° to 77°F)
- protect from excessive moisture

Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate

Questions or comments?

1-800-CLARITIN (1-800-252-7484) or **www.claritin.com**



Dist by: Bayer HealthCare LLC
Whippany, NJ 07981
Product of Ireland.

Bayer

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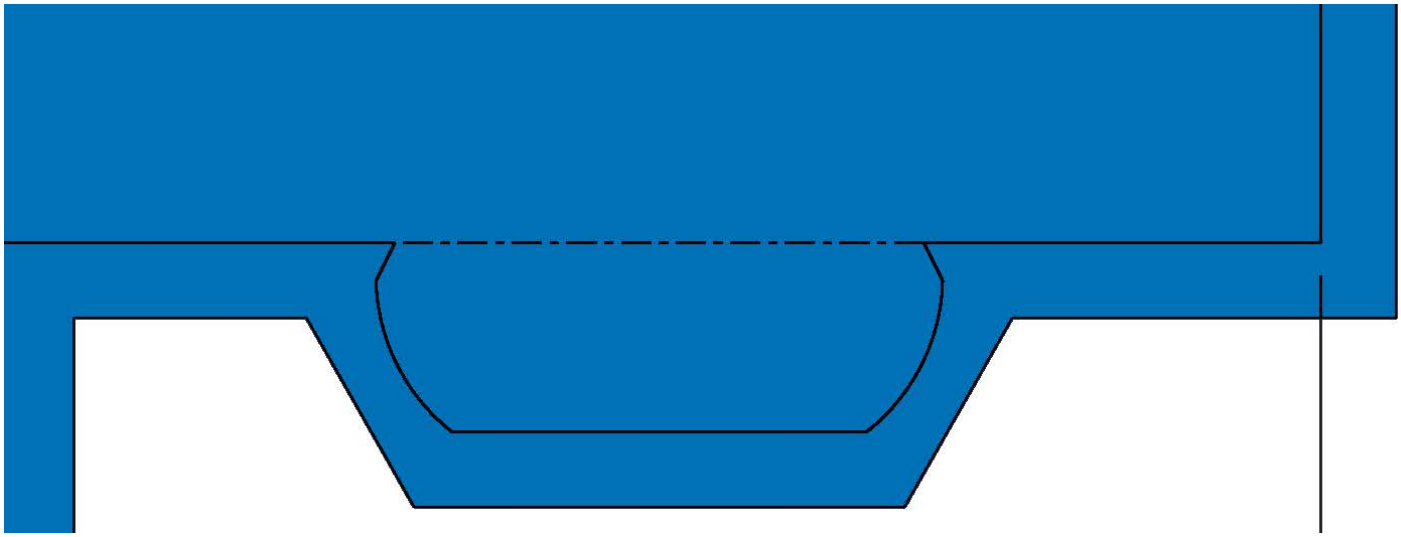

CLARITIN

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-7237
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics			
Color	white	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	458;Claritin10
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11523-7237-1	1 in 1 CARTON	04/12/1993	
1		45 in 1 BOTTLE; Type 0: Not a Combination		

1		Product		
2	NDC:11523-7237-3	7 in 1 CARTON	08/12/1993	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11523-7237-5	1 in 1 BLISTER PACK	04/12/1993	
3		90 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:11523-7237-6	3 in 1 CARTON	04/12/1993	
4		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:11523-7237-7	4 in 1 CARTON	04/12/1993	
5		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:11523-7237-8	1 in 1 CARTON	04/12/1993	
6		108 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:11523-7237-9	1 in 1 CARTON	10/05/2017	
7		90 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019658	04/12/1993	

CLARITIN

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-7160
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics			
Color	white	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	458;Claritin10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-7160-1	1 in 1 CARTON	04/12/1993	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11523-7160-2	1 in 1 CARTON	04/12/1993	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11523-7160-3	2 in 1 CARTON	04/12/1993	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:11523-7160-5	3 in 1 CARTON	04/12/1993	
4		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:11523-7160-6	1 in 1 CARTON	04/12/1993	
5		2 in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:11523-7160-7	1 in 1 POUCH	04/12/1993	
6		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
7	NDC:11523-7160-9	1 in 1 CARTON	04/12/1993	
7		60 in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:11523-7160-8	1 in 1 CARTON	09/01/2016	
8		60 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019658	04/12/1993	

CLARITIN

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-1527
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics			
Color	white	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	458;Claritin10
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11523-1527-1	50 in 1 BOX	04/12/1993	
1		1 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:11523-1527-2	50 in 1 BOX	04/12/1993	
2		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019658	04/12/1993	

CLARITIN

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-4359
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	458;Claritin10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-4359-1	1 in 1 CARTON	04/12/1993	
1		40 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11523-4359-2	1 in 1 CARTON	04/12/1993	
2		100 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:11523-4359-3	1 in 1 CARTON	11/01/2017	
3		90 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:11523-4359-4	1 in 1 CARTON	02/15/2019	
4		55 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:11523-4359-5	1 in 1 CARTON	02/15/2019	
5		85 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:11523-4359-6	1 in 1 CARTON	12/01/2019	
6		110 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:11523-4359-8	2 in 1 BLISTER PACK	04/12/1993	
7		60 in 1 BOTTLE; Type 0: Not a Combination		

		Product		
8	NDC:11523-4359-9	1 in 1 CARTON	12/01/2019	
8		35 in 1 BOTTLE; Type 0: Not a Combination Product		
9	NDC:11523-4359-7	1 in 1 BLISTER PACK	04/12/1993	
9		115 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019658	04/12/1993	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 10/2022 Bayer HealthCare LLC.